# **PCT**

# WORLD INTELLECTUAL PROPERTY ORGANIZATION International Bureau



## INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(51) International Patent Classification <sup>5</sup> :		(11) International Publication Number	WO 91/19528
A61M 29/00, 25/01	A1	(43) International Publication Date:	26 December 1991 (26.12.91)

(21) International Application Number:

PCT/US91/03509

(22) International Filing Date:

17 May 1991 (17.05.91)

(30) Priority data:

535,932

11 June 1990 (11.06.90) US

(71) Applicant: SCHNEIDER (USA) INC. [US/US]; 5905 Nathan Lane, Plymouth, MN 55442 (US).

(72) Inventor: SHOCKEY, Rick, L.; 3890 Westbury Drive, Eagan, MN 55123 (US).

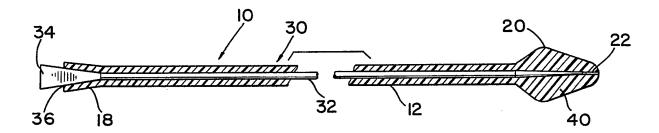
(74) Agents: RICHARDSON, Peter, C. et al.; Pfizer Inc., 235 East 42nd Street, New York, NY 10017 (US).

(81) Designated States: AT (European patent), AU, BE (European patent), CA, CH (European patent), DE, DE (Utility model), DE (European patent), DK (European patent), ES (European patent), FR (European patent), GB (European patent), GR (European patent), IT (European patent), JP, LU (European patent), NL (European patent), SE (European patent).

#### **Published**

With international search report.

(54) Title: TRACKING GUIDEWIRE



#### (57) Abstract

A guidewire and guidewire assembly (10) for placement within a blood vessel for penetrating an occlusion therein. The guidewire comprises a length of flexible wire (12) having a concentric lumen (14) running its entire length and a distal end portion (20) having an arcuate tip (22) and a diameter greater than that of the wire immediately proximal thereto. The guidewire assembly comprises the guidewire described along with a flexible stylet (32) substantially the same length as the flexible wire and disposed within the lumen of the wire. In operation, the distal end portion is positioned in the blood vessel against an occlusion, and a dottering action is thereafter provided whereby the distal end portion repeatedly impinges on the occlusion until penetration of the occlusion occurs.

## FOR THE PURPOSES OF INFORMATION ONLY

Codes used to identify States party to the PCT on the front pages of pamphlets publishing international applications under the PCT.

AT	Austria	ES	Spain	MG	Madagascar
ΑÜ	Australia	FI	Finland	ML	Mali
BB	Barbados	FR	France	MN	Mongolia
BE	Belgium	GA	Gabon	MR	Mauritania
BF	Burkina Faso	GB	United Kingdom	MW	Malawi
BG	Bulgaria	GN	Guinea	NL	Netherlands
BJ	Benin	GR	Greece	NO	Norway
BR	Brazil	HU	Hungary	PL	Poland
CA	Canada	IT	Italy	RO	Romania
CF	Central African Republic	JP	Japan	SD	Sudan
CG	Congo	KP	Democratic People's Republic	SE	Sweden
CH	Switzerland		of Korea	SN	Senegal
Ci	Côte d'Ivoire	KR	Republic of Korea	SU	Soviet Union
CM	Cameroon	LI	Licchtenstein	TD	Chad
CS	Czechoslovakia	LK	Sri Lanka	TG	Togo
DE	Germany	LU	Luxembourg	US	United States of America
DK	Denmark	MC	Monaco		

PCT/US91/03509 WO 91/19528

-1-

# TRACKING GUIDEWIRE Technical Field

This invention relates generally to a tracking guidewire, and in particular to an occlusion-penetrable guidewire having a lumen throughout its entire length into which a stylet can be inserted, and having a distal end portion with an arcuate tip and a diameter greater than that of the immediately proximal wire.

### Background Art

10

25

Vessel entry for treatment of certain untoward health conditions is a common practice. Such entry can include insertion into a blood vessel of a guidewire whose distal end is expected to reach a certain site within the body and have utility thereafter as 15 required. Many times, however, a blood vessel may be completely or almost completely occluded, thereby rendering it substantially impossible to advance a guidewire there beyond to a designated site without first employing a separate procedure to remove the 20 occlusion.

It is therefore a primary object of the present invention to provide a guidewire having a lumen running its entire length and a distal end portion capable of penetrating a vascular occlusion. Another object of the present invention is to provide such a guidewire wherein the tip of the distal end portion is arcuate and the diameter of the distal end portion is greater than that of the immediately proximal wire. another object of the present invention is to provide a guidewire assembly wherein a stylet can be removably inserted into the length of the guidewire lumen to thereby enhance guidewire structure. These and other objects will become apparent throughout the description which now follows.

-2-

#### Disclosure of Invention

The present invention is a guidewire for placement within a blood vessel for penetrating an occlusion in the vessel. The guidewire comprises a length of 5 flexible wire having a concentric lumen running its entire length. At its proximal end the wire has a hub opening to the lumen, and at its distal end it has a distal end portion which has an arcuate tip and a diameter greater than the diameter of the wire 10 immediately proximal thereto. The invention additionally includes a guidewire assembly which comprises the guidewire and a flexible substantially the length of the wire for removable insertion into the lumen of the wire for substantially 15 its entire length. Such stylet placement provides a greater stiffness and structural integrity to the guidewire.

Finally a method of penetrating an occlusion in a blood vessel comprises inserting the guidewire or guidewire assembly into an occluded blood vessel and positioning the arcuate tip of the distal end portion against the proximal wall of the occlusion. After such positioning, a dottering action is provided to the guidewire or guidewire assembly for a sufficient period of time whereby the repeated impingement upon the occlusion results in penetration thereof. Once the distal portion of the guidewire or guidewire assembly has passed through the occlusion, and the stylet has been withdrawn in the case of the guidewire assembly, a contrast medium can be injected into the lumen to confirm guidewire positioning within the vessel.

#### Brief Description of Drawings

Presently preferred embodiments of the invention are illustrated in the accompanying drawings in which:

5

25

30

Figure 1 is a side elevation view of a guidewire;
Figure 2 is a side elevation view partially in section of a guidewire assembly;

Figure 3 is a side elevation view of a stylet;
Figure 4 is a side elevation view of a second embodiment of a distal end of a guidewire; and

Figure 5 is a side elevation view of a third embodiment of a distal end of a guidewire.

## Modes for Carrying Out the Invention

guidewire 10 Referring to Figure 1, a 1.0 illustrated. The guidewire 10 comprises a length of flexible wire 12 having a concentric lumen 14 running the entire length of the wire 12 and a proximal end hub 18 which opens to the lumen 14. The opposite end of the wire 12 has a distal end portion 20 which has an 15 arcuate tip 22 and a diameter greater than the diameter of the wire 12 immediately proximal to the end portion 20. The guidewire 10 can be conventionally constructed of metal core, and is preferably constructed of metal coils. 20

Figure 2 illustrates a guidewire assembly 30 comprising a guidewire 10, as shown in Figure 1, and a flexible stylet 32, as shown in Figure 3, disposed within the lumen 14 of the wire 12. The stylet 32 is wire 12 and length of the substantially the additionally has a proximal end member 34 whose shape is complimentary to the interior wall 36 of the hub 18. The hub 18 is provided with locking means such as conventional Luer locking threads which securely maintains the stylet 32 within the lumen 14, yet provides releasability for withdrawal of the stylet 32 as required. The stylet 32 can be constructed of metal or polymer core, and is preferably constructed of metal.

-4-

As earlier noted, the distal end portion 20 of the wire 12 has an arcuate tip 22 and has a diameter which is greater than that of the wire 12 immediately proximal thereto. Three non-limiting examples of 5 preferred shapes of the end portion 20 are illustrated in Figures 1, 2, 4 and 5. Specifically, Figures 1 and 2 show an arrow shape 40; Figure 4 shows an elliptical shape 42; and Figure 5 illustrates a tear-drop shape It is to be understood, of course, that shapes 10 other than those illustrated can be employed as long as an arcuate tip is provided to thereby enhance physical intrusion of an occlusion. The various shapes of the respective distal end portions are attained in the manufacturing process which can include EDM machining 15 and grinding.

In operation, the guidewire 10, and in particular its distal arcuate tip 22 and distal end portion 20, functions to penetrate an occlusion in a blood vessel. The user inserts the guidewire 10 into a blood vessel 20 and positions the arcuate tip 22 against the proximal wall of an occlusion. Once placed, the guidewire 10 is subjected to a dottering action by the user to effectuate a repeated impinging action upon the occlusion by the distal end portion 20 for a period of 25 time sufficient to penetrate the entire length of the occlusion and thereby permit continued travel of the guidewire 10 itself or of other apparatus. occlusion exhibit a resistance too great to permit the guidewire 10 alone to penetrate, the stylet 32 30 inserted into the length of the lumen 14 and locked into place. The inserted stylet 32 provides a greater stiffness and structure to the guidewire 10, thereby providing a more aggressive impingement action on the occlusion during dottering to accomplish penetration

-5-

thereof. Once the distal end portion 20 of the guidewire 10 is through the stenosis, the stylet 32 can be removed from the lumen 14 and a contrast medium can injected into the lumen 14 to thereby confirm the true position of the distal end portion 20.

While an illustrative and presently preferred embodiment of the invention has been described in detail herein, it is to be understood that the inventive concepts may be otherwise variously embodied and employed and that the appended claims are intended to be construed to include such variations except insofar as limited by the prior art.

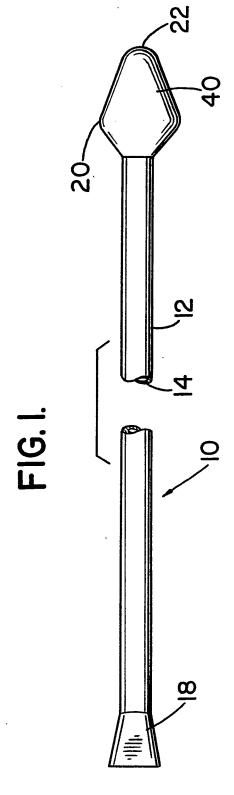
#### CLAIMS

1) A guidewire assembly (10) for placement within a blood vessel for penetrating an occlusion therein, the guidewire assembly (10) comprising:

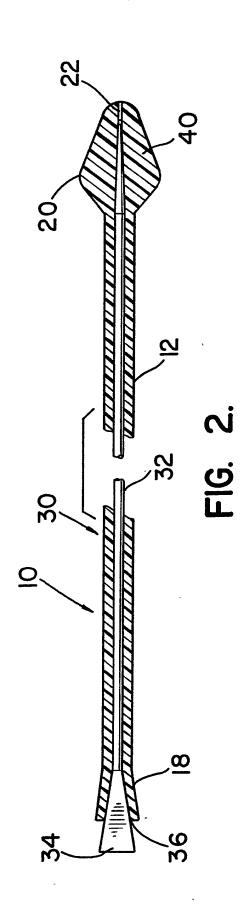
- a) a length of flexible wire (12) having a concentric lumen (14) running the entire length of the wire (12), with said wire (12) having a proximal end hub (18) opening to the lumen (14) and a distal end portion (20) having an arcuate tip (22) and a diameter greater than the diameter of the wire (14) immediately proximal thereto; and
- b) a flexible stylet (32) substantially the length of the wire (12) and additionally having a proximal end member (34) complimentarily shaped to the interior wall (36) of the hub (18), said stylet (32) having a diameter less than the diameter of the lumen (14) of the wire (12) and removably insertable within the lumen (14) substantially throughout the entire length of said lumen (14).
- 2) A guidewire assembly (10) as claimed in claim 1 and having in addition locking means disposed at the proximal end hub (18) opening of the wire (12) for releasably securing the proximal end member (34) of the stylet (32) within at least a portion of the hub (18).
- 25 3) A guidewire assembly (10) as claimed in claim 2 wherein the distal end portion (20) of the wire (12) is in the shape of an arrow (40).
- 4) A guidewire assembly (10) as claimed in claim 2 wherein the distal end portion (20) of the wire (12) 30 is in the shape of an ellipse (42).
  - 5) A guidewire assembly (10) as claimed in claim 2 wherein the distal end portion (20) of the wire (12) is in the shape of a tear drop (44).
    - 6) A guidewire assembly (10) as claimed in claim

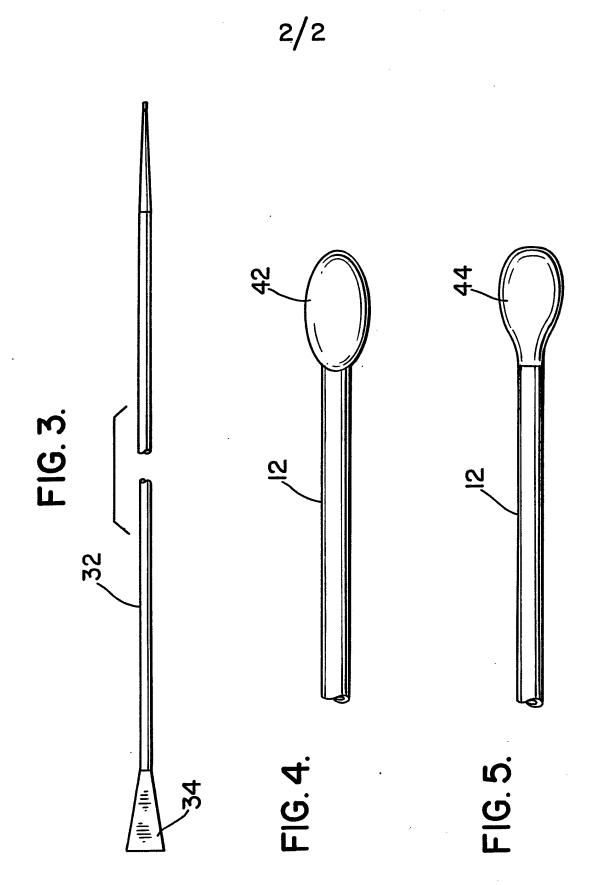
- 1 wherein the distal end portion (20) of the wire (12) is in the shape of an arrow (40).
- 7) A guidewire assembly (10) as claimed in claim 1 wherein the distal end portion (20) of the wire (12) is in the shape of an ellipse (42).
  - 8) A guidewire assembly (10) as claimed in claim 1 wherein the distal end portion (20) of the wire (12) is in the shape of a tear drop (44).

1/2









**SUBSTITUTE SHEET** 

			International Application No	
I. CLASSI	FICATION OF SUBJ	ECT MATTER (if several classification sy	ymbols apply, indicate all) <sup>6</sup>	
According	to International Paten	t Classification (IPC) or to both National Cl		
Int.	C1. 5	A61M29/00; A61M25/01		·
II. FIELDS	SEARCHED			
	•	Minimum Docume	entation Searched <sup>7</sup>	
Classificat	tion System		Classification Symbols	
Int.	C1. 5	A61M; A61B		
	· · · · · · · · · · · · · · · · · · ·	Documentation Searched other to the Extent that such Documents a	than Minimum Documentation ure Included in the Fields Searched <sup>8</sup>	
III. DOCU		D TO BE RELEVANT <sup>9</sup>		The State of the S
Category °	Citation of Do	ocument, 11 with indication, where appropria	ite, of the relevant passages 12	Relevant to Claim No. <sup>13</sup>
		(UTLLED M ) 11.	07 1065	1-2,4-5,
Y	US,A,3 .	196 876 (MILLER M.) Jul	y 2/, 1965	1-2,4-5, 7-8
	see cla	im; figures 1-4		, 5
	366 610	III, FIGURES A 7		
Υ	US,A,4 388 076 (WATERS) June 14, 1983			1-2,4-5, 7-8
	see abstract; figures 1-5			
A	EP.A.363	3 661 (ADVANCED CARDIOVA	ASCULAR SYSTEMS	1-2
	INC.) A	oril 18, 1990		
	see absi	tract; figures 2-4		
	see colu	umn 3, line 52 - line 54	4	
٨	ED A 2 3	290 917 (LINDEMANN) June	e 11 1976	3-8
A	rk, M, Z Z	e 4, line 19 - line 29;	figures 2-3	
	see page			
Α	US,A,3 9	999 551 (SPITZ ET AL.) [	December 28, 1976	3,6
	see colu	umn 3, line 36 - line 47	7; figures 2,4	
		on as as		
° Special	categories of cited doc	currents: 10	"T" later document published after the internat	ional filing date
"A" doc	ument defining the gen	eral state of the art which is not	or priority date and not in conflict with the cited to understand the principle or theory	underlying the
	sidered to be of particulier document but publi	nar relevance shed on or after the international	invention  "X" document of particular relevance; the claim	sed invention
fili	ng date	v doubts on priority claim(s) or	cannot be considered novel or cannot be co involve an inventive step	ensid <b>ered</b> to
whi	ch is cited to establish tion or other special re	the publication date of another	"Y" document of particular relevance; the claim cannot be considered to involve an inventiv	ed invention
"O" doc	ument referring to an o	oral disclosure, use, exhibition or	document is combined with one or more of	her such docu-
other means  "P" document published prior to the international filing date but  "P" document published prior to the international filing date but				
late	r than the priority date	daimed	"&" document member of the same patent fami	ly
IV. CERTII	FICATION			
Date of the	Actual Completion of the	he International Search	Date of Mailing of this International Searc	h Report
	30 J	IULY 1991	13, 89, 91	,
			Signature of Authorized Officer	
International	Searching Authority		MIR Y GUILLEN V.	Mrs 1
	EUROPEA	N PATENT OFFICE	MIK I GOTELLIA A.	/ -

EUROPEAN PATENT OFFICE

### ANNEX TO THE INTERNATIONAL SEARCH REPORT ON INTERNATIONAL PATENT APPLICATION NO.

PCT/US 91/03509

47937

This annex lists the patent family members relating to the patent documents cited in the above-mentioned international search report. The members are as contained in the European Patent Office EDP file on

The European Patent Office is in no way liable for these particulars which are merely given for the purpose of information.

30/0

30/07/91

Patent document cited in search report	Publication date	Patent family member(s)		Publication date
US-A-3196876		None		
US-A-4388076	14-06-83	US-E-	RE32306	16-12-86
EP-A-363661	18-04-90	US-A- JP-A-	4917102 2191467	17-04-90 27-07-90
FR-A-2290917	11-06-76	DE-A- CH-A- US-A-	2454351 591254 4013079	20-05-76 15-09-77 22-03-77
US-A-3999551	28-12-76	None		